''(A) have been approved or indexed under the relevant provision of the Public Health Service Act or Federal Food, Drug, and Cosmetic Act; and

''(B) have permission for commercial marketing or use.

''(2) In this subsection, the term 'covered date' means the later of—

''(A) the date an application is approved—

''(i) under section 351(a)(2)(C) of the Public Health Service Act; or

''(ii) under section 505(b) or 512(c) of the Federal Food, Drug, and Cosmetic Act;

''(B) the date an application is conditionally approved under section 571(b) of the Federal Food, Drug, and Cosmetic Act;

''(C) the date a request for indexing is granted under section 572(d) of the Federal Food, Drug, and Cosmetic Act; or

''(D) the date of issuance of the interim final rule controlling the drug under section 201(j) of the Controlled Substances Act.''.

SEC. 3. ENHANCING NEW DRUG DEVELOPMENT.

Section 303 of the Controlled Substances Act (21 U.S.C. 823) is amended by adding at the end the following:

''(i)(1) For purposes of registration to manufacture a controlled substance under subsection (d) for use only in a clinical trial, the Attorney General shall register the applicant, or serve an order to show cause upon the applicant in accordance with section 304(c), not later than 180 days after the date on which the application is accepted for filing.

''(2) For purposes of registration to manufacture a controlled substance under subsection (a) for use only in a clinical trial, the Attorney General shall, in accordance with the regulations issued by the Attorney General, issue a notice of application not later than 90 days after the application is accepted for filing. Not later than 90 days after the date on which the period for comment pursuant to such notice ends, the Attorney General shall register the applicant, or serve an order to show cause upon the applicant in accordance with section 304(c), unless the Attorney General has granted a hearing on the application under section 1008(i) of the Controlled Substances Import and Export Act.''.

SEC. 4. RE-EXPORTATION AMONG MEMBERS OF THE EUROPEAN ECONOMIC AREA.

Section 1003 of the Controlled Substances Import and Export Act (21 U.S.C. 953) is amended—

(1) in subsection(f)—

(A) in paragraph (5)—

(i) by striking ''(5)'' and inserting ''(5)(A)'';

(ii) by inserting '', except that the controlled substance may be exported from a second country that is a member of the European Economic Area to another country that is a member of the European Economic Area, provided that the first country is also a member of the European Economic Area'' before the period at the end; and

(iii) by adding at the end the following:

''(B) Subsequent to any re-exportation described in subparagraph (A), a controlled substance may continue to be exported from any country that is a member of the European Economic Area to any other such country, if—

''(i) the conditions applicable with respect to the first country under paragraphs (1), (2), (3), (4), (6), and (7) are met by each subsequent country from which the controlled substance is exported pursuant to this paragraph; and

''(ii) the conditions applicable with respect to the second country under paragraphs (1), (2), (3), (4), (6), and (7) are met by each subsequent country to which the controlled substance is exported pursuant to this paragraph.''; and

(B) in paragraph (6)—

(i) by striking ''(6)'' and inserting ''(6)(A)''; and

(ii) by adding at the end the following:

''(B) In the case of re-exportation among members of the European Economic Area, within 30 days after each re-exportation, the person who exported the controlled substance from the United States delivers to the Attorney General—

''(i) documentation certifying that such re-exportation has occurred; and

''(ii) information concerning the consignee, country, and product.''; and

(2) by adding at the end the following:

''(g) LIMITATION.—Subject to paragraphs (5) and (6) of subsection (f) in the case of any controlled substance in schedule I or II or any narcotic drug in schedule III or IV, the Attorney General shall not promulgate nor enforce any regulation, subregulatory guidance, or enforcement policy which impedes re-exportation of any controlled substance among European Economic Area countries, including by promulgating or enforcing any requirement that—

''(1) re-exportation from the first country to the second country or re-exportation from the second country to another country occur within a specified period of time; or

''(2) information concerning the consignee, country, and product be provided prior to exportation of the controlled substance from the United States or prior to each re-exportation among members of the European Economic Area.''.

The PRESIDING OFFICER. The Senator from Ohio.

WOUNDED WARRIORS FEDERAL LEAVE ACT OF 2015

Mr. PORTMAN. Madam President, I ask unanimous consent that the Committee on Homeland Security and Governmental Affairs be discharged from further consideration of H.R. 313 and the Senate proceed to its immediate consideration.

The PRESIDING OFFICER. Without objection, it is so ordered.

The clerk will report the bill by title.

The senior assistant legislative clerk read as follows:

A bill (H.R. 313) to amend title 5, United States Code, to provide leave to any new Federal employee who is a veteran with a service-connected disability rated at 30 percent or more for purposes of undergoing medical treatment for such disability, and for other purposes.

There being no objection, the Senate proceeded to consider the bill.

Mr. PORTMAN. I ask unanimous consent that the bill be read a third time and passed and the motion to reconsider be considered made and laid upon the table.

The PRESIDING OFFICER. Without objection, it is so ordered.

The bill (H.R. 313) was ordered to a third reading, was read the third time, and passed.

IMPROVING REGULATORY TRANSPARENCY FOR NEW MEDICAL THERAPIES ACT

Mr. PORTMAN. Madam President, I ask unanimous consent that the Committee on Health, Education, Labor, and Pensions be discharged from further consideration of H.R. 639 and the Senate proceed to its immediate consideration.

The PRESIDING OFFICER. Without objection, it is so ordered.

The clerk will report the bill by title.

The senior assistant legislative clerk read as follows:

A bill (H.R. 639) to amend the Controlled Substances Act with respect to drug scheduling recommendations by the Secretary of Health and Human Services, and with respect to registration of manufacturers and distributors seeking to conduct clinical testing.

There being no objection, the Senate proceeded to consider the bill.

Mr. PORTMAN. I ask unanimous consent that the substitute amendment, which is at the desk, be considered and agreed to, the bill, as amended, be read a third time and passed, and the motion to reconsider be considered made and laid upon the table.

The PRESIDING OFFICER. Without objection, it is so ordered.

The amendment (No. 2748) in the nature of a substitute was agreed to.

(The amendment is printed in today's RECORD under ''Text of Amendments.'')

The amendment was ordered to be engrossed, and the bill to be read a third time.

The bill was read the third time.

The bill (H.R. 639), as amended, was passed.

CONGRATULATING THE MINNESOTA LYNX ON THEIR VICTORY IN THE 2015 WOMEN'S NATIONAL BASKETBALL ASSOCIATION FINALS

Mr. PORTMAN. Madam President, I ask unanimous consent that the Senate proceed to the immediate consideration of S. Res. 297, submitted earlier today.

The PRESIDING OFFICER. The clerk will report the resolution by title.

The senior assistant legislative clerk read as follows:

A resolution (S. Res. 297) congratulating the Minnesota Lynx on their victory in the 2015 Women's National Basketball Association Finals.

There being no objection, the Senate proceeded to consider the resolution.

Mr. PORTMAN. Madam President, I ask unanimous consent that the resolution be agreed to, the preamble be agreed to, and the motions to reconsider be considered made and laid upon the table with no intervening action or debate.

The PRESIDING OFFICER. Without objection, it is so ordered.

The resolution (S. Res. 297) was agreed to.

The preamble was agreed to.

(The resolution, with its preamble, is printed in today's RECORD under ''Submitted Resolutions.'')

ORDERS FOR TUESDAY, OCTOBER 27, 2015

Mr. PORTMAN. Madam President, I ask unanimous consent that when the